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EXAMINER

FORMAN, BETTY J

ART UNIT	PAPER NUMBER
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1634

DATE MAILED: 12/11/2003

Please find below and/or attached an Office communication concerning this application or proceeding.

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Office Action Summary

Application No.

09/636,387

Applicant(s)

STUELPNAGEL ET AL.

Examiner

BJ Forman

Art Unit

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-- The MAILING DATE of this communication appears on the cover sheet with the correspondence address --
Period for Reply

A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) FROM THE MAILING DATE OF THIS COMMUNICATION.

- Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.
- If the period for reply specified above is less than thirty (30) days, a reply within the statutory minimum of thirty (30) days will be considered timely.
- If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.
- Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133).
- Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).

Status

- 1) ☒ Responsive to communication(s) filed on 30 September 2003.
- 2a) ☐ This action is **FINAL**. 2b) ☒ This action is non-final.
- 3) ☐ Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under *Ex parte Quayle*, 1935 C.D. 11, 453 O.G. 213.

Disposition of Claims

- 4) ☒ Claim(s) 49-54 and 61-75 is/are pending in the application.
- 4a) Of the above claim(s) _____ is/are withdrawn from consideration.
- 5) ☐ Claim(s) _____ is/are allowed.
- 6) ☒ Claim(s) 49-54 and 61-75 is/are rejected.
- 7) ☐ Claim(s) _____ is/are objected to.
- 8) ☐ Claim(s) _____ are subject to restriction and/or election requirement.

Application Papers

- 9) ☒ The specification is objected to by the Examiner.
- 10) ☐ The drawing(s) filed on _____ is/are: a) ☐ accepted or b) ☐ objected to by the Examiner.
 Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).
 Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).
- 11) ☐ The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.

Priority under 35 U.S.C. §§ 119 and 120

- 12) ☐ Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).
 a) ☐ All b) ☐ Some * c) ☐ None of:
 1. ☐ Certified copies of the priority documents have been received.
 2. ☐ Certified copies of the priority documents have been received in Application No. _____.
 3. ☐ Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).
 * See the attached detailed Office action for a list of the certified copies not received.
- 13) ☒ Acknowledgment is made of a claim for domestic priority under 35 U.S.C. § 119(e) (to a provisional application) since a specific reference was included in the first sentence of the specification or in an Application Data Sheet. 37 CFR 1.78.
 a) ☐ The translation of the foreign language provisional application has been received.
- 14) ☒ Acknowledgment is made of a claim for domestic priority under 35 U.S.C. §§ 120 and/or 121 since a specific reference was included in the first sentence of the specification or in an Application Data Sheet. 37 CFR 1.78.

Attachment(s)

- 1) ☐ Notice of References Cited (PTO-892) 4) ☐ Interview Summary (PTO-413) Paper No(s). _____
- 2) ☐ Notice of Draftsperson's Patent Drawing Review (PTO-948) 5) ☐ Notice of Informal Patent Application (PTO-152)
- 3) ☐ Information Disclosure Statement(s) (PTO-1449) Paper No(s) _____ 6) ☐ Other: _____

DETAILED ACTION

Continued Examination Under 37 CFR 1.114

1. A request for continued examination under 37 CFR 1.114, including the fee set forth in 37 CFR 1.17(e), was filed in this application after final rejection. Since this application is eligible for continued examination under 37 CFR 1.114, and the fee set forth in 37 CFR 1.17(e) has been timely paid, the finality of the previous Office action has been withdrawn pursuant to 37 CFR 1.114. Applicant's submission filed on 30 September 2003 has been entered.

Status of the Claims

2. This action is in response to papers filed 30 September 2003 in which Applicant presented argument regarding the previous rejections. The previous rejections in the Office Action dated 30 May 2003 are maintained. Applicant's arguments have been considered and are addressed below.

The "Listing of Claims" submitted on 30 September 2003, lists Claims 1-48 and 55-60 as "withdrawn". These claims were cancelled in papers filed 25 June 2001. Therefore, Applicant's Listing of Claims is incorrect.

Claims 49-54 and 61-75 are pending and under prosecution.

Specification

3. The amendment filed 25 June 2001 is objected to under 35 U.S.C. 132 because it introduces new matter into the disclosure. 35 U.S.C. 132 states that no amendment shall introduce new matter into the disclosure of the invention. The added material which is not supported by the original disclosure is as follows: The amendment adds language to the first paragraph of the specification to incorporate by reference parent applications 09/500,555 and 60/119,323. Because the parent applications were not incorporated by reference in the originally filed specification, the amendment constitutes new matter.

Applicant is required to cancel the new matter in the reply to this Office Action.

Claim Rejections - 35 USC § 103

4. The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negated by the manner in which the invention was made.

5. Claims 49-54, 61-63, 65-69 and 71-74 are rejected under 35 U.S.C. 103(a) as being unpatentable over Walt et al. (U.S. Patent No. 6,327,410 B1, filed 11 September 1998) in view of Brenner (U.S. Patent No. 5,863,722, filed 7 June 1995) and in view of the definitions of Morris ed. (Academic Press Dictionary of Science and Technology, Academic Press, 1992, page 821).

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Regarding Claim 49, Walt et al. teach a method of determining the presence of a target analyte in a sample comprising: acquiring a first data image of a random array composition comprising: a substrate with a surface comprising discrete sites; a population of microspheres comprising at least a first and a second sub-population each comprising a bioactive agent; and a fiducial (i.e. marker bead, Column 19, lines 2-5) wherein said microspheres are randomly distributed on said surface such that said discrete sites contain microspheres; using the fiducial to register said first data image to create a registered first data image; contacting said random array with said sample; acquiring a second data image from said array with said sample; using the fiducial to register said second data image to create a registered second data image; and comparing said first and said second registered data images to determine the presence or absence of said target analyte (Column 18, line 59-Column 19, line 62 and Claims 17-21) wherein additional, non-optical signature encoding, is used e.g. size, shape and/or dimension of the microspheres and/or optical fibers which suggests that a population does not have an optical signature (Column 19, lines 6-30) but they do not specifically teach at least one microsphere subpopulation does not have an optical signature. However, microspheres without optical signatures were well known in the art at the time the claimed invention was made as taught by Brenner et al. who teach a similar array composition comprising: a substrate with a surface comprising discrete sites; and a population of microspheres comprising at least a first and second subpopulation wherein each subpopulation comprises a bioactive agent wherein the microspheres are distributed on said surface wherein the microsphere does not comprise an optical signature and wherein microspheres size and shape are selected based on experimental design (Column 9, lines 62-65; Column 19, line 20-49; and Column 21, line 15-Column 22, line 61). It would have been obvious to one of ordinary skill in the art at the time the claimed invention was made to apply the microspheres not having an optical signature as taught by Brenner et al and the non-optical signature encoding taught by Walt et al to the microsphere compositions of Walt et al and to provide at least one

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subpopulation of microsphere without an optical signature thereby eliminating the need to provide optical signatures on all the microspheres for the obvious benefits of simplicity. One skilled in the art would have been further motivated to modify the microspheres of Walt et al by replacing the optical signature microspheres of one subpopulation with non-labeled microspheres to be used to detect labeled targets wherein only microspheres bound to labeled targets are detected for the obvious benefits of target-specific detection.

Furthermore, in the example provided the array is excited to provide a registered image (Fig. 10A), a sample is added, the array is again excited to provide a second registered image (Fig. 10B). The first and second images are compared to determine the presence of the target analyte (antibody, Column 27, lines 30-54). In this embodiment, Walt does not specifically teach utilizing the fiducial to create the images. However, they teach that their fiducials are an important element in identifying beads on the array (Column 19, lines 1-5) which clearly suggests that they utilize the fiducial to create the data image. Therefore, It would have been obvious to one of ordinary skill in the art at the time the claimed invention was made to utilize the fiducials of Walt et al to create their images for the expected benefit of identifying beads one-from-another as they desire (Column 19, lines 1-5).

Regarding Claim 50, Walt et al teach the method wherein the array comprises a fiber optic bundle and the registration of the first data image utilizes a fiducial fiber (Column 19, lines 2-53).

Regarding Claim 51, Walt et al teach the method wherein the array comprises microspheres and the registration of the first data image utilizes a fiducial microsphere i.e. marker bead (Column 19, lines 2-53).

Regarding Claim 52, Walt et al teach the method wherein the registration utilizes a fiducial template i.e. subarray bundle (Column 18, lines 59-66).

Regarding Claim 53, Walt et al. teach the method wherein the bioactive agents are proteins (Column 7, lines 55-61).

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Regarding Claim 54, Walt et al. teach the method wherein the bioactive agents are nucleic acids (Column 7, lines 55-61).

Regarding Claim 61, Walt et al. teach a method of determining the presence of a target analyte in a sample comprising: providing a first data image of a random array comprising: a substrate with a surface comprising discrete sites; a population of microspheres comprising at least a first and a second sub-population each comprising a bioactive agent; and a fiducial (i.e. marker bead, Column 19, lines 2-5) wherein said microspheres are randomly distributed on said surface such that said discrete sites contain microspheres; contacting said random array with said sample; acquiring a second data image from said array with said sample; using the fiducial to register said second data image to create a registered second data image; and comparing said first and said second registered data images to determine the presence or absence of said target analyte (Column 18, line 59-Column 19, line 62; Example 1; and Claims 17-21) wherein additional, non-optical signature encoding, is used e.g. size, shape and/or dimension of the microspheres and/or optical fibers (Column 19, lines 6-30) but they do not specifically teach at least one microsphere subpopulation does not have an optical signature.

However, microspheres without optical signatures were well known in the art at the time the claimed invention was made as taught by Brenner et al. who teach a similar array composition comprising: a substrate with a surface comprising discrete sites; and a population of microspheres comprising at least a first and second subpopulation wherein each subpopulation comprises a bioactive agent wherein the microspheres are distributed on said surface wherein the microsphere does not comprise an optical signature and wherein microspheres size and shape are selected based on experimental design (Column 9, lines 62-65; Column 19, line 20-49; and Column 21, line 15-Column 22, line 61). It would have been obvious to one of ordinary skill in the art at the time the claimed invention was made to apply the microspheres not having an optical signature as taught by Brenner et al and the non-optical signature encoding taught by Walt et al to the microsphere compositions of Walt et al

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and to provide at least one subpopulation of microsphere without an optical signature thereby eliminating the need to provide optical signatures on all the microspheres for the obvious benefits of simplicity. One skilled in the art would have been further motivated to modify the microspheres of Walt et al by replacing the optical signature microspheres of one subpopulation with non-labeled microspheres to be used to detect labeled targets wherein only microspheres bound to labeled targets are detected for the obvious benefits of target-specific detection.

Regarding Claim 62, Walt et al. teach the method of Claim 49 wherein the substrate is selected from the group consisting of glass and plastic i.e. optical fibers (Column 5, lines 57-60) which are comprised of glass or plastic as defined by Morris ed. (page 821).

Regarding Claim 63, Walt et al teach the method wherein the registration utilizes a fiducial edge i.e. subarray bundle (Column 18, lines 59-66).

Regarding Claim 65, Walt et al. teach the method of Claims 53 and 54 wherein the substrate is selected from the group consisting of glass and plastic i.e. optical fibers (Column 5, lines 57-60) which are comprised of glass or plastic as defined by Morris ed. (page 821).

Regarding Claim 66, Walt et al. teach the method of Claims 49 and 62 wherein each sub-population comprises a unique optical signature (Column 13, lines 8-24).

Regarding Claim 67, Walt et al. teach the method wherein the said unique optical signature is a bleed-through signature i.e. the signal is obtained from multiple wavelengths (Column 14, line 17-67).

Regarding Claim 68, Walt et al. teach the method of Claims 49 and 62 wherein each sub-population comprises an identifier binding ligand that will bind a decoder binding ligand such that the identification of the bioactive agent can be elucidated i.e. enzyme and substrate whereby the enzyme is identified in the presence of the substrate (Column 20, line 51-Column 21, line 6).

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Regarding Claim 69, Walt et al teach the array comprises at least three fiducial and each is a fiducial fiber i.e. a unique tag for each of 100 different subarrays (Column 18, line 59-Column 19, line 5).

Regarding Claim 71, Walt et al teach the array wherein the fiducial fibers have a different color i.e. unique tag (Column 18, lines 59-66).

Regarding Claim 72, Walt et al teach the array wherein the registration utilizes at least three fiducial and each of said fiducials is a microsphere i.e. marker beads which identify each of 100 subarrays (Column 18, line 57-column 19, line 5).

Regarding Claim 73, Walt et al teach the fiducials have different sizes (Column 19, lines 6-30).

Regarding Claim 74, Walt et al teach the array wherein the fiducial fibers have a different color i.e. unique tag (Column 18, lines 59-66).

Response to Arguments

6. Applicant argues that neither Brenner et al nor Walt et al teach or suggest using a fiducial to create a registered first and second data image.

Applicant acknowledges that Walt et al teaches using fiducials for positional coding, but argues that the positional coding of Walt et al can be determined without creating a registered data image as instantly claimed.

The argument has been considered but is not found persuasive because Applicant has not provided factual evidence for their assertion that the positional coding of Walt et al can be determined without creating a registered data image.

The arguments of counsel cannot take the place of evidence in the record. In re Schulze, 346 F.2d 600, 602, 145 USPQ 716, 718 (CCPA 1965). Examples of attorney statements which are not evidence and which must be supported by an appropriate affidavit or declaration include statements regarding unexpected results, commercial success, solution of a long-felt need, inoperability of the prior art, invention before the date of the reference, and allegations that the author(s) of the prior art derived the disclosed subject matter from the applicant. (see (MPEP 716.01(c)).

Furthermore, the cited passage of Walt et al is concerned with positional coding utilizing fiducials (marker beads/bead size/bead shape) to identify one bundle from another (Column 19, lines 23) wherein additional coding parameters include sequential loading of the bead populations randomly onto the array followed by reading an optical signature from each sequentially loaded population to provide a matrices of bead location (Column 19, lines 31-52).

In the example provided by Walt, the array is excited to provide a registered image (Fig. 10A), a sample is added, the array is again excited to provide a second registered image (Fig. 10B). The first and second images are compared to determine the presence of the target analyte (antibody, Column 27, lines 30-54).

The claims are drawn to "using" a fiducial. The claims do not limit or define how the fiducial is used in the method. As such, any use of the fiducial in the creation of an image meets the limitations of the claims. Walt et al clearly use the fiducials to make, encode and decode the array (Column 18, line 59-Column 19, line 62 and Claims 17-21). Therefore, Walt et al teaches the use as claimed.

Applicant argues that the references do not teach or suggest a motivation to combine their teachings and further argue that the mere fact that references can be combined does not render the combination obvious. In response to applicant's argument that there is no suggestion to combine the references, the examiner recognizes that obviousness can only be established by combining or modifying the teachings of the prior art to produce the claimed invention where there is some teaching, suggestion, or motivation to do so found either in the references themselves or in the knowledge generally available to one of ordinary skill in the art. See *In re Fine*, 837 F.2d 1071, 5 USPQ2d 1596 (Fed. Cir. 1988) and *In re Jones*, 958 F.2d 347, 21 USPQ2d 1941 (Fed. Cir. 1992). In this case, one of ordinary skill in the art would have been motivated provide at least one subpopulation of not having an optical signature to thereby eliminating the need to provide optical signatures on all the microspheres for the obvious

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benefits of simplicity. The courts have stated that it would be obvious to omit an element when a function attributed to said element is not desired or required (see *Ex parte Wu*, 10 USPQ 2031). Brenner et al teach the optical signature is not required for target detection because the target is detected via the tag (Column 19, lines 20-49). It would have been obvious to one of ordinary skill in the art at the time the claimed invention was made to eliminate the optical signature of Walt et al based on that fact that the optical signature is not required and therefore for the obvious benefits of simplicity. One skilled in the art would have been further motivated to modify the microspheres of Walt et al by replacing the optical signature microspheres of one subpopulation with non-labeled microspheres to be used to detect labeled targets wherein only microspheres bound to labeled targets are detected for the obvious benefits of target-specific detection.

7. Claims 64, 70 and 75 are rejected under 35 U.S.C. 103(a) as being unpatentable over Walt et al. (U.S. Patent No. 6,327,410, filed 11 September 1998) in view of Brenner (U.S. Patent No. 5,863,722, filed 7 June 1995) and in view of the definitions of Morris ed. (Academic Press Dictionary of Science and Technology, Academic Press, 1992, page 821) as applied to Claims 49 and 61 above and further in view of Augenlicht (U.S. Patent No. 4,981,783, filed 16 April 1986)

Regarding Claim 64, Walt et al. teach a method of determining the presence of a target analyte in a sample comprising: acquiring a first data image of a random array composition comprising: a substrate with a surface comprising discrete sites; a population of microspheres comprising at least a first and a second sub-population each comprising a bioactive agent; and a fiducial (i.e. marker bead, Column 19, lines 2-5) wherein said microspheres are randomly distributed on said surface such that said discrete sites contain microspheres; using the

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fiducial to register said first data image to create a registered first data image; contacting said random array with said sample; acquiring a second data image from said array with said sample; using the fiducial to register said second data image to create a registered second data image; and comparing said first and said second registered data images to determine the presence or absence of said target analyte (Column 18, line 59-Column 19, line 62 and Claims 17-21) wherein additional, non-optical signature encoding, is used e.g. size, shape and/or dimension of the microspheres and/or optical fibers (Column 19, lines 6-30) and Brenner et al. who teach a similar array composition comprising microspheres which do not comprise an optical signature and wherein microspheres size and shape are selected based on experimental design (Column 9, lines 62-65; Column 19, line 20-49; and Column 21, line 15-Column 22, line 61). One skilled in the art would have been motivated to modify the microspheres of Walt et al by replacing the optical signature microspheres of one subpopulation with non-labeled microspheres as taught by Brenner et al to thereby eliminate the need to provide optical signatures on all the microspheres for the obvious benefits of simplicity. Walt et al do not teach at least a first edge of the array is a fiducial edge. However, Augenlicht teaches a similar method for determining the presence of a target comprising a substrate with a surface comprising discrete sites and at least a first and second population each comprising a bioactive agent distributed on the surface; acquiring a data image to create a registered image and comparing registered images to determine the presence of said target wherein the registered image utilizes a fiducial marking (Column 7, lines 18-46) wherein at least a first edge of the array is a fiducial edge i.e. upper right edge, upper left edge and lower left edge (Fig. 1) and wherein the fiducial markings permit rapid and accurate automated scanning to thereby identify targets rapidly and accurately (Column 8, lines 15-29). It would have been obvious to one of ordinary skill in the art at the time the claimed invention was made to modify the substrate of Walt et al. by utilizing a fiducial edge as suggested by Augenlicht to thereby reduce operator time and human error by permitting automated scanning of the substrate for the

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expected benefit of rapid and accurate target analysis as suggested by Augenlicht (Column 8, lines 15-29).

Regarding Claim 70, Walt et al teach the array comprises at least three fiducial and each is a fiducial fiber i.e. a unique tag for each of 100 different subarrays (Column 18, line 59-Column 19, line 5) wherein the fiducials are beads of different sizes (Column 19, lines 6-30) and they further teach that the beads are not uniform, but are instead irregularly shaped (Column 7, lines 32-40) which suggest that the bead have different shapes. Furthermore, the skilled practitioner in the art would have been further motivated to modify the different sized fiducial markers of Walt et al by providing at least one fiducial having a different shape to thereby obtain a substrate having fiducials of differing and identifiable shape (e.g. a different identifiable shape in each corner) for the obvious benefit of facilitating identification the target analyte. Because subarrays each having a fiducial of different shape would permit identification of the subarray by simple visual identification of fiducial shape thereby facilitating identification of the subarray and target analyte. Therefore, it would have been obvious to one of ordinary skill in the art at the time the claimed invention was made to modify the fiducial beads of Walt et al by encompassing fiducials of different (irregular) shaped into each subarray for the obvious benefits of facilitating target analyte identification.

Regarding Claim 75, the array wherein the registration utilizes at least three fiducial and each of said fiducials is a microsphere i.e. marker beads which identify each of 100 subarrays (Column 18, line 57-column 19, line 5) and they also teach the fiducials are identified by size (Column 19, lines 7-12) and therefore the fiducials are not identified by label which suggests that the fiducials are not labeled. However, Walt et al do not specifically teach unlabeled fiducials. It would have been obvious to one of ordinary skill in the art at the time the claimed invention was made to modify the different-sized fiducials of Walt et al by providing unlabeled fiducials of different size based on the suggestion of Walt et al to thereby eliminate the labeling step for the obvious benefit of convenience and economy of time and labor.

Response to Arguments

8. Applicant argues that Augenlicht does not cure the deficiencies of Walt et al and Brenner et al. The argument has been considered but is not found persuasive for the reasons stated above.

Applicant argues that the cited references do not teach or suggest a motivation to combine or modify their teachings. In response to applicant's argument that there is no suggestion to combine the references, the examiner recognizes that obviousness can only be established by combining or modifying the teachings of the prior art to produce the claimed invention where there is some teaching, suggestion, or motivation to do so found either in the references themselves or in the knowledge generally available to one of ordinary skill in the art. See *In re Fine*, 837 F.2d 1071, 5 USPQ2d 1596 (Fed. Cir. 1988) and *In re Jones*, 958 F.2d 347, 21 USPQ2d 1941 (Fed. Cir. 1992). In this case, one of ordinary skill in the art would have been motivated provide at least one subpopulation of not having an optical signature to thereby eliminating the need to provide optical signatures on all the microspheres for the obvious benefits of simplicity. The courts have stated that it would be obvious to omit an element when a function attributed to said element is not desired or required (see *Ex parte Wu*, 10 USPQ 2031). Brenner et al teach the optical signature is not required for target detection because the target is detected via the tag (Column 19, lines 20-49). It would have been obvious to one of ordinary skill in the art at the time the claimed invention was made to eliminate the optical signature of Walt et al based on that fact that the optical signature is not required and therefore for the obvious benefits of simplicity. One skilled in the art would have been further motivated to modify the microspheres of Walt et al by replacing the optical signature microspheres of one subpopulation with non-labeled microspheres to be used to detect labeled targets wherein only microspheres bound to labeled targets are detected for the obvious benefits of target-specific detection. Furthermore, It would have been obvious to one of ordinary skill in the art the fiducials of Walt et al. by utilizing a fiducial edge as suggested by

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Augenlicht to thereby reduce operator time and human error by permitting automated scanning of the substrate for the expected benefit of rapid and accurate target analysis as suggested by Augenlicht (Column 8, lines 15-29).

Applicant argues that it is unclear how the triangular fiducials of Augenlicht are placed on their array or how they could be applied to the array of Walt. Hence, Applicant argues, Augenlicht fails to enable one of ordinary skill to make the array of Walt et al having the fiducials of Augenlicht. The argument has been considered but is not found persuasive because as cited above, Walt et al clearly suggest the fiducials (marker beads) have different shape because they teach that they are "irregular" (Column 7, lines 32-40).

In response to applicant's argument that the references fail to show certain features of applicant's invention, it is noted that the features upon which applicant relies (i.e., triangular fiducials) are not recited in the rejected claim(s). Although the claims are interpreted in light of the specification, limitations from the specification are not read into the claims. See *In re Van Geuns*, 988 F.2d 1181, 26 USPQ2d 1057 (Fed. Cir. 1993).

Conclusion

9. No claim is allowed.

10. Any inquiry concerning this communication or earlier communications from the examiner should be directed to BJ Forman whose telephone number is (703) 306-5878 until 13 January 2004. Starting 14 January 2004, the examiner's phone number will be (517) 272-0741. The examiner can normally be reached on 6:00 TO 3:30 Monday through Thursday and alternate Fridays.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Gary Benzion can be reached on (703) 308-1119. The fax phone numbers for the organization where this

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application or proceeding is assigned are (703) 872-9306 for regular communications and (703) 308-8724 for After Final communications.

Any inquiry of a general nature or relating to the status of this application or proceeding should be directed to the receptionist whose telephone number is (703) 308-0196.



BJ Forman, Ph.D.
Primary Examiner
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December 5, 2003